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(54) Title: MUTANT LUCIFERASES (57) Abstract <p>The invention provides active, non-naturally occurring mutants of beetle luciferases and DNAs which encode such mutants. A mutant luciferase of the invention differs from the corresponding wild-type luciferase by producing bioluminescence with a wavelength of peak intensity that differs by at least 1 nm from the wavelength of peak intensity of the bioluminescence produced by the wild-type enzyme. The mutant luciferases and DNAs of the invention are employed in various biosensing applications.</p>			

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MUTANT LUCIFERASES

TECHNICAL FIELD

This invention generally relates to luciferase enzymes that produce luminescence, like that from fireflies. More particularly, the invention concerns mutant luciferases of beetles. The mutant luciferases of the invention are made by genetic engineering, do not occur in nature, and, in each case, include modifications which cause a change in color in the luminescence that is produced. The luciferases of the invention can be used, like their naturally occurring counterparts, to provide luminescent signals in tests or assays for various substances or phenomena.

BACKGROUND OF THE INVENTION

The use of reporter molecules or labels to qualitatively or quantitatively monitor molecular events is well established. They are found in assays for medical diagnosis, for the detection of toxins and other substances in industrial environments, and for basic and applied research in biology, biomedicine, and biochemistry. Such assays include immunoassays, nucleic acid probe hybridization assays, and assays in which a reporter enzyme or other protein is produced by expression under control of a particular promoter. Reporter molecules, or labels in such assay systems, have included radioactive isotopes, fluorescent agents, enzymes and chemiluminescent agents.

Included in the assay system employing chemiluminescence to monitor or measure events of interest are assays which measure the activity of a bioluminescent enzyme, luciferase.

Light-emitting systems have been known and isolated from many luminescent organisms including bacteria, protozoa, coelenterates, molluscs, fish, millipedes, flies, fungi, worms, crustaceans, and beetles, particularly click beetles of genus *Pyrophorus* and the fireflies of the genera *Photinus*, *Photuris*, and

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Luciola. In many of these organisms, enzymes catalyze monooxygenations and utilize the resulting free energy to excite a molecule to a high energy state. Visible light is emitted when the excited molecule spontaneously
5 returns to the ground state. This emitted light is called "bioluminescence." Hereinafter it may also be referred to simply as "luminescence."

The limited occurrence of natural bioluminescence is an advantage of using luciferase enzymes as reporter
10 groups to monitor molecular events. Because natural bioluminescence is so rare, it is unlikely that light production from other biological processes will obscure the activity of a luciferase introduced into a biological system. Therefore, even in a complex environment, light
15 detection will provide a clear indication of luciferase activity.

Luciferases possess additional features which render them particularly useful as reporter molecules for biosensing (using a reporter system to reveal properties
20 of a biological system). Signal transduction in biosensors (sensors which comprise a biological component) generally involves a two step process: signal generation through a biological component, and signal transduction and amplification through an electrical component.
25 Signal generation is typically achieved through binding or catalysis. Conversion of these biochemical events into an electrical signal is typically based on electrochemical or caloric detection methods, which are limited by the free energy change of the biochemical
30 reactions. For most reactions this is less than the energy of hydrolysis for two molecules of ATP, or about 70 kJ/mole. However, the luminescence elicited by luciferases carries a much higher energy content. Photons emitted from the reaction catalyzed by firefly
35 luciferase (560 nm) have 214 KJ/einstein. Furthermore, the reaction catalyzed by luciferase is one of the most efficient bioluminescent reactions known, having a

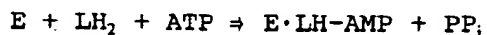
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quantum yield of nearly 0.9. This enzyme is therefore an extremely efficient transducer of chemical energy.

Since the earliest studies, beetle luciferases, particularly that from the common North American firefly species *Photinus pyralis*, have served as paradigms for understanding of bioluminescence. The fundamental knowledge and applications of luciferase have been based on a single enzyme, called "firefly luciferase," derived from *Photinus pyralis*. However, there are roughly 1800 species of luminous beetles worldwide. Thus, the luciferase of *Photinus pyralis* is a single example of a large and diverse group of beetle luciferases. It is known that all beetle luciferases catalyze a reaction of the same substrate, a polyheterocyclic organic acid, D-(-)-2-(6'-hydroxy-2'-benzothiazolyl)- Δ^2 -thiazoline-4-carboxylic acid (hereinafter referred to as "luciferin", unless otherwise indicated), which is converted to a high energy molecule. It is likely that the catalyzed reaction entails the same mechanism in each case.

The general scheme involved in the mechanism of beetle bioluminescence appears to be one by which the production of light takes place after the oxidative decarboxylation of the luciferin, through interaction of the oxidized luciferin with the enzyme. The color of the light apparently is determined by the spatial organization of the enzyme's amino acids which interact with the oxidized luciferin.

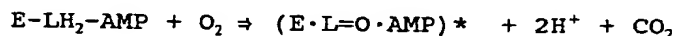
The luciferase-catalyzed reaction which yields bioluminescence (hereinafter referred to simply as "the luciferase-luciferin reaction") has been described as a two-step process involving luciferin, adenosine triphosphate (ATP), and molecular oxygen. In the initial reaction, the luciferin and ATP react to form luciferyl adenylate with the elimination of inorganic pyrophosphate, as indicated in the following reaction:



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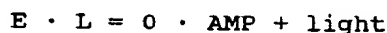
where E is the luciferase, LH_2 is luciferin, and PPi is pyrophosphate. The luciferyl adenylate, $\text{LH}_2\text{-AMP}$, remains tightly bound to the catalytic site of luciferase. When this form of the enzyme is exposed to molecular oxygen, the enzyme-bound luciferyl adenylate is oxidized to yield oxyluciferin ($\text{L}=\text{O}$) in an electronically excited state. The excited oxidized luciferin emits light on returning to the ground state as indicated in the following reaction:

10



↓

15



One quantum of light is emitted for each molecule of luciferin oxidized. The electronically excited state of the oxidized luciferin is a characteristic state of the luciferase-luciferin reaction of a beetle luciferase; the color (and, therefore, the energy) of the light emitted upon return of the oxidized luciferin to the ground state is determined by the enzyme, as evidenced by the fact that various species of beetles having the same luciferin emit differently colored light.

Luciferases have been isolated directly from various sources. The cDNAs encoding luciferases of various beetle species have been reported. (See de Wet et al., *Molec. Cell. Biol.* 7, 725 - 737 (1987); Masuda et al., *Gene* 77, 265 - 270 (1989); Wood et al., *Science* 244, 700 - 702 (1989)). With the cDNA encoding a beetle luciferase in hand, it is entirely straightforward for the skilled to prepare large amounts of the luciferase by isolation from bacteria (e.g., *E. coli*), yeast, mammalian cells in culture, or the like, which have been transformed to express the cDNA. Alternatively, the cDNA, under control of an appropriate promoter and other

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signals for controlling expression, can be used in such a cell to provide luciferase, and ultimately bioluminescence catalyzed thereby, as a signal to indicate activity of the promoter. The activity of the promoter may, in turn, reflect another factor that is sought to be monitored, such as the concentration of a substance that induces or represses the activity of the promoter. Various cell-free systems, that have recently become available to make proteins from nucleic acids encoding them, can also be used to make beetle luciferases.

Further, the availability of cDNAs encoding beetle luciferases and the ability to rapidly screen for cDNAs that encode enzymes which catalyze the luciferase-luciferin reaction (see de Wet et al., supra and Wood et al., supra) also allow the skilled to prepare, and obtain in large amounts, other luciferases that retain activity in catalyzing production of bioluminescence through the luciferase-luciferin reaction. These other luciferases can also be prepared, and the cDNAs that encode them can also be used, as indicated in the previous paragraph. In the present disclosure, the term "beetle luciferase" or "luciferase" means an enzyme that is capable of catalyzing the oxidation of luciferin to yield bioluminescence, as outlined above.

The ready availability of cDNAs encoding beetle luciferases makes possible the use of the luciferases as reporters in assays employed to signal, monitor or measure genetic events associated with transcription and translation, by coupling expression of such a cDNA, and consequently production of the enzyme, to such genetic events.

Firefly luciferase has been widely used to detect promoter activity in eucaryotes. Though this enzyme has also been used in procaryotes, the utility of firefly luciferase as genetic reporter in bacteria is not commonly recognized. As genetic reporters, beetle

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luciferases are particularly useful since they are monomeric products of a single gene. In addition, no post-translational modifications are required for enzymatic activity, and the enzyme contains no prosthetic groups, bound cofactors, or disulfide bonds.

5 Luminescence from *E.coli* containing the gene for firefly luciferase can be triggered by adding the substrate luciferin to the growth medium. Luciferin readily penetrates biological membranes and cannot be used as a

10 carbon or nitrogen source by *E.coli*. The other substrates required for the bioluminescent reaction, oxygen and ATP, are available within living cells. However, measurable variations in luminescence color from luciferases would be needed for systems which utilize two

15 or more different luciferases as reporters (signal generators).

Clones of different beetle luciferases, particularly of a single genus or species, can be utilized together in bioluminescent reporter systems.

20 Expression in exogenous hosts should differ little between these luciferases because of their close sequence similarity. Thus, in particular, the click beetle luciferases may provide a multiple reporter system that can allow the activity of two or more different promoters

25 to be monitored within a single host, or for different populations of cells to be observed simultaneously. The ability to distinguish each of the luciferases in a mixture, however, is limited by the width of their emissions spectra.

30 One of the most spectacular examples of luminescence color variation occurs in *Pyrophorus plagiophthalmus*, a large click beetle indigenous to the Caribbean. This beetle has two sets of light organs, a pair on the dorsal surface of the prothorax, and a single

35 organ in a ventral cleft of the abdomen. Four different luciferase clones have been isolated from the ventral organ. The luciferin-luciferase reactions catalyzed by

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these enzymes produces light that ranges from green to orange.

Spectral data from the luciferase-luciferin reaction catalyzed by these four luciferases show four overlapping peaks of nearly even spacing, emitting green (peak intensity: 546 nanometers), yellow-green (peak intensity: 560 nanometers), yellow (peak intensity: 578 nanometers) and orange (peak intensity: 593 nanometers) light. The respective proteins are named LucPpLGR, LucPpLYG, LucPpLYE and LucPpLOR. Though the wavelengths of peak intensity of the light emitted by these luciferases range over nearly 50 nm, there is still considerable overlap among the spectra, even those with peaks at 546 and 593 nm. Increasing the difference in wavelength of peak intensity would thus be useful to obtain greater measurement precision in systems using two or more luciferases.

The amino acid sequences of the four luciferases from the ventral organ are highly similar. Comparisons of the sequences show them to be 95 to 99% identical.

It would be desirable to enhance the utility of beetle luciferases for use in systems using multiple reporters to effect mutations in luciferase-encoding cDNAs to produce mutant luciferases which, in the luciferase-luciferin reaction, produce light with differences between wavelengths of peak intensity that are greater than those available using currently available luciferases.

Beetle luciferases are particularly suited for producing these mutant luciferases since color variation is a direct result of changes in the amino acid sequence.

Mutant luciferases of fireflies of genus *Luciola* are known in the art. Kajiyama et al., U.S. Patent Nos. 5,219,737 and 5,229,285.

In using luciferase expression in eukaryotic cells for biosensing, it would be desirable to reduce transport of the luciferase to peroxisomes. Sommer et

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al., Mol. Biol. Cell 3, 749 - 759 (1992), have described mutations in the three carboxy-terminal amino acids of *P. pyralis* luciferase that significantly reduce peroxisome-targeting of the enzyme.

- 5 The sequences of cDNAs encoding various beetle luciferases, and the amino acid sequences deduced from the cDNA sequences, are known, as indicated in Table I.

Table I

10 References for cDNA and Amino Acid Sequences of Various Wild-Type Beetle Luciferases

	Luciferase	Reference
15	LucPp1GR	K. Wood, Ph.D. Dissertation, University of California, San Diego (1989), see also SEQ ID NO:1; Wood et al., Science 244, 700-702 (1989)
20	LucPp1YG	K. Wood, Ph.D. Dissertation, University of California, San Diego (1989); Wood et al., Science 244, 700-702 (1989)
25	LucPp1YE	K. Wood, Ph.D. Dissertation, University of California, San Diego (1989); Wood et al., Science 244, 700-702 (1989)
30	LucPp1OR	K. Wood, Ph.D. Dissertation, University of California, San Diego (1989); Wood et al., Science 244, 700-702 (1989)
35	Photinus pyralis	de Wet et al., Mol. Cell. Biol. 7, 725 - 737 (1987); K. Wood, Ph.D. Dissertation, University of California, San Diego (1989); Wood et al., Science 244, 700 - 702 (1989)
40	Luciola cruciata	Kajiyama et al., United States Patent No. 5,229,285; Masuda et al., United States Patent No. 4,968,613
45	Luciola lateralis	Kajiyama et al., United States Patent No. 5,229,285
50	Luciola mingrelica	Devine et al., Biochim. et Biophys. Acta 1173, 121-132(1993)

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The cDNA and amino acid sequences of LucPplGR, the green-emitting luciferase of the elaterid beetle *Pyrophorus plagiophthalmus*, are shown in SEQ ID NO:1.

5 SUMMARY OF THE INVENTION

The present invention provides mutant luciferases of beetles and DNAs which encode the mutant luciferases. Preferably, the mutant luciferases produce a light of different color from that of the corresponding wild-type
10 luciferase and preferably this difference in color is such that the wavelength of peak intensity of the luminescence of the mutant differs by at least 1 nm from that of the wild-type enzyme.

The mutant luciferases of the invention differ
15 from the corresponding wild-type enzymes by one or more, but typically fewer than three, amino acid substitutions. The luciferases of the invention may also entail changes in one or more of the three carboxy-terminal amino acids to reduce peroxisome targeting.

20 In one surprising aspect of the invention, it has been discovered that combining in a single mutant two amino acid substitutions, each of which, by itself, occasions a change in color (shift in wavelength of peak intensity) of bioluminescence, causes the mutant to have
25 a shift in wavelength of peak intensity that is greater than either shift caused by the single amino acid substitutions.

cDNAs encoding the mutant luciferases of the invention may be obtained straightforwardly by any
30 standard, site-directed mutagenesis procedure carried out with a cDNA encoding the corresponding wild-type enzyme or another mutant. The mutant luciferases of the invention can be made by standard procedures for expressing the cDNAs which encode them in prokaryotic or
35 eukaryotic cells.

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A fuller appreciation of the invention will be gained upon examination of the following detailed description of the invention.

5 DETAILED DESCRIPTION OF THE INVENTION

In the following description and examples, process steps are carried out and concentrations are measured at room temperature (about 20 °C to 25 °C) and atmospheric pressure unless otherwise specified.

10 All amino acids referred to in the specification, except the non-enantiomorphous glycine, are L-amino acids unless specified otherwise. An amino acid may be referred to using the one-letter or three-letter designation, as indicated in the following Table II.

15 Table II

Designations for Amino Acids

Amino Acid	Three-Letter Designation	One-Letter Designation
20 L-alanine	Ala	A
L-arginine	Arg	R
L-asparagine	Asn	N
L-aspartic acid	Asp	D
L-cysteine	Cys	C
25 L-glutamic acid	Glu	E
L-glutamine	Gln	Q
glycine	Gly	G
L-histidine	His	H
L-isoleucine	Ile	I
30 L-leucine	Leu	L
L-lysine	Lys	K
L-methionine	Met	M
L-phenylalanine	Phe	F
L-proline	Pro	P
35 L-serine	Ser	S
L-threonine	Thr	T
L-tryptophan	Trp	W
L-tyrosine	Tyr	Y
L-valine	Val	V

40 "X" means any one of the twenty amino acids listed in Table II.

Peptide or polypeptide sequences are written and numbered from the initiating methionine, which is numbered "1," to the carboxy-terminal amino acid.

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A substitution at a position in a polypeptide is indicated with [designation for original amino acid]_[position number][designation for replacing amino acid]. For example, substitution of an alanine at position 100 in a polypeptide with a glutamic acid would be indicated by Ala₁₀₀Glu or A₁₀₀E. Typically, the substitution will be preceded by a designation for the polypeptide in which the substitution occurs. For example, if the substitution A₁₀₀E occurs in an hypothetical protein designated "Luck," the substitution would be indicated as Luck-Ala₁₀₀Glu or Luck-A₁₀₀E. If there is more than one substitution in a polypeptide, the indications of the substitutions are separated by slashes. For example, if the hypothetical protein "Luck" has a substitution of glutamic acid for alanine at position 100 and a substitution of asparagine for lysine at position 150, the polypeptide with the substitutions would be indicated as Luck-Ala₁₀₀Glu/Lys₁₅₀Asn or Luck-A₁₀₀E/K₁₅₀N. To indicate different substitutions at a position in a polypeptide, the designations for the substituting amino acids are separated by commas. For example, if the hypothetical "Luck" has substitutions of glutamic acid, glycine or lysine for alanine at position 100, the designation would be Luck-Ala₁₀₀/Glu,Gly,Lys or Luck-A₁₀₀/E,G,K.

The standard, one-letter codes "A," "C," "G," and "T" are used herein for the nucleotides adenylate, cytidylate, guanylate, and thymidylate, respectively. The skilled will understand that, in DNAs, the nucleotides are 2'-deoxyribonucleotide-5'-phosphates (or, at the 5'-end, triphosphates) while, in RNAs, the nucleotides are ribonucleotide-5'-phosphates (or, at the 5'-end, triphosphates) and uridylate (U) occurs in place of T. "N" means any one of the four nucleotides.

Oligonucleotide or polynucleotide sequences are written from the 5'-end to the 3'-end.

The term "mutant luciferase" is used herein to refer to a luciferase which is not naturally occurring

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and has an amino acid sequence that differs from those of naturally occurring luciferases.

In one of its aspects, the present invention is a mutant beetle luciferase which produces bioluminescence (i.e., catalyzes the oxidation of luciferin to produce bioluminescence) which has a shift in wavelength of peak intensity of at least 1 nm from the wavelength of peak intensity of the bioluminescence produced by the corresponding wild-type luciferase and has an amino acid sequence that differs from that of the corresponding wild-type luciferase by a substitution at one position or substitutions at two positions; provided that, if there is a substitution at one position, the position corresponds to a position in the amino acid sequence of LucPplGR selected from the group consisting of position 214, 215, 223, 224, 232, 236, 237, 238, 242, 244, 245, 247, 248, 282, 283 and 348; provided further that, if there are substitutions at two positions, at least one of the positions corresponds to a position in the amino acid sequence of LucPplGR selected from the group consisting of position 214, 215, 223, 224, 232, 236, 237, 238, 242, 244, 245, 247, 248, 282, 283 and 348; and provided that the mutant optionally has a peroxisome-targeting-avoiding sequence at its carboxy-terminus.

Exemplary mutant luciferases of the invention are those of the group consisting of LucPplGR-R₂₁₅H, -R₂₁₅G, -R₂₁₅T, -R₂₁₅M, -R₂₁₅P, -R₂₁₅A, -R₂₁₅L, -R₂₂₃L, -R₂₂₃Q, -R₂₂₃M, -R₂₂₃H, -V₂₂₄I, -V₂₂₄S, -V₂₂₄F, -V₂₂₄Y, -V₂₂₄L, -V₂₂₄H, -V₂₂₄G, -V₂₃₂E, -V₂₃₆H, -V₂₃₆W, -Y₂₃₇S, -Y₂₃₇C, -L₂₃₈R, -L₂₃₈M, -L₂₃₈Q, -L₂₃₈S, -L₂₃₈D, -H₂₄₂A, -F₂₄₄L, -G₂₄₅S, -G₂₄₅E, -S₂₄₇H, -S₂₄₇T, -S₂₄₇Y, -S₂₄₇F, -I₂₄₈R, -I₂₄₈V, -I₂₄₈F, -I₂₄₈T, -I₂₄₈S, -I₂₄₈N, -H₃₄₈N, -H₃₄₈Q, -H₃₄₈E, -H₃₄₈C, -S₂₄₇F/F₂₄₆L, -S₂₄₇F/I₂₄₈C, -S₂₄₇F/I₂₄₈T, -V₂₂₄F/R₂₁₅G, -V₂₂₄F/R₂₁₅T, -V₂₂₄F/R₂₁₅V, -V₂₂₄F/R₂₁₅P, -V₂₂₄F/P₂₂₂S, -V₂₂₄F/Q₂₂₇E, -V₂₂₄F/L₂₃₈V, -V₂₂₄F/L₂₃₈T, -V₂₂₄F/S₂₄₇G, -V₂₂₄F/S₂₄₇H, -V₂₂₄F/S₂₄₇T, and -V₂₂₄F/S₂₄₇F.

The following Table III shows spectral properties of these and other exemplary mutant luciferases.

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TABLE III

Protein		Spectral Properties		
	LucPplGR-	peak	shift	width
5	w.t.	545	0	72
	V ₂₁₄ S	*		
	Q	*		
	Y	*		
	K	*		
10	L	*		
	G	*		
	C	*		
	E	*		
	F	*		
15	P	*		
	H	*		
	R	*		
	R ₂₁₃ H	562	17	82
	Q	567	22	81
20	G	576	31	82
	T	576	31	84
	M	582	37	83
	P	588	43	91
	S	*		
25	Y	*		
	K	*		
	L	*		
	C	*		
	E	*		
30	F	*		
	R ₂₂₃ L	549	4	75
	Q	549	4	73

*Spectral shift (≥ 2 nm) observed by eye.

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TABLE III, cont.

Protein		Spectral Shift		
	LucPplGR-	peak	shift	width
5	R ₂₂₃ M	549	4	75
	H	551	6	75
	S	*		
	Y	*		
	K	*		
10	G	*		
	C	*		
	E	*		
	F	*		
	P	*		
15	V ₂₂₄ I	546	1	75
	S	556	11	70
	F	561	16	84
	Y	565	20	87
	L	578	33	94
20	H	584	39	69
	G	584	39	70
	V ₂₃₂ E	554	9	83
	V ₂₃₆ H	554	9	74
	W	554	9	74
25	Y ₂₃₇ S	553	8	73
	C	554	9	74
	L ₂₃₈ R	544	-1	72
	M	555	10	75
	Q	557	12	76
30	S	559	14	73
	D	568	23	76
	H ₂₄₂ A	559	14	75

*Spectral shift (≥ 2 nm) observed by eye.

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TABLE III, cont.

Protein	Spectral Properties		
LucPplGR-	peak	shift	width
H ₂₄₂ S	561	16	74
F ₂₄₄ L	555	10	73
G ₂₄₅ S	558	13	75
E	574	29	79
S ₂₄₇ H	564	19	72
Y	566	21	79
F	569	24	84
I ₂₄₈ R	544	-1	72
V	546	1	72
F	548	3	74
T	554	9	75
S	558	13	80
N	577	32	90
H ₃₄₈ A	592	47	67
C	593	48	66
N	597	52	67
Q	605	60	72
V ₂₁₄ C/V ₂₂₄ A	559	14	72
S ₂₄₇ F/F ₂₄₆ L	567	22	79
S ₂₄₇ F/I ₂₄₈ C	586	41	84
S ₂₄₇ F/I ₂₄₈ T	596	51	80
T ₂₃₃ A/L ₂₃₈ M	555	10	75
V ₂₈₂ I/I ₂₈₃ V	563	3	73
V ₂₂₄ F/R ₂₁₅ G	584	39	80
V ₂₂₄ F/R ₂₁₅ T	587	42	80
V ₂₂₄ F/R ₂₁₅ V	589	44	80
V ₂₂₄ F/R ₂₁₅ P	597	52	81
V ₂₂₄ F/P ₂₂₂ S	564	3	86

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TABLE III, cont.

Protein	Spectral Properties		
LucPplGR-	peak	shift	width
V ₂₂₄ F/Q ₂₂₇ E	583	38	85
V ₂₂₄ F/L ₂₃₈ V	575	30	85
V ₂₂₄ F/L ₂₃₈ M	576	31	87
V ₂₂₄ F/S ₂₄₇ G	581	36	84
V ₂₂₄ F/S ₂₄₇ H	581	36	79
V ₂₂₄ F/S ₂₄₇ Y	595	50	88
V ₂₂₄ F/S ₂₄₇ F	597	52	85

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"Corresponding positions" in luciferases other than LucPplGR can be determined either from alignments at the amino acid level that are already known in the art (see, e.g., Wood et al., Science 244, 700 - 702 (1989);
5 Devine et al., Biochim. et Biophys. Acta 1173, 121-132(1993)) or by simply aligning at the amino acid level to maximize alignment of identical or conservatively substituted residues, and keeping in mind in particular that amino acids 195 - 205 in the LucPplGR sequence are
10 very highly conserved in all beetle luciferases and that there are no gaps for more than 300 positions after that highly conserved 11-mer in any beetle luciferase amino acid sequence.

A "peroxisome-targeting-avoiding sequence at its
15 carboxy-terminus" means (1) the three carboxy-terminal amino acids of the corresponding wild-type luciferase are entirely missing from the mutant; or (2) the three carboxy-terminal amino acids of the corresponding wild-type luciferase are replaced with a sequence, of one, two
20 or three amino acids that, in accordance with Sommer et al., supra, will reduce peroxisome-targeting by at least 50 %. If the three carboxy-terminal amino acids of the wild-type luciferase are replaced by a three-amino-acid peroxisome-targeting-avoiding sequence in the mutant, and
25 if the sequence in the mutant is $X_1X_2X_3$, where X_3 is carboxy-terminal, then X_1 is any of the twenty amino acids except A, C, G, H, N, P, Q, T and S, X_2 is any of the twenty amino acids except H, M, N, Q, R, S and K, and X_3 is any of the twenty amino acids except I, M, Y and L.
30 Further, any one or two, or all three, of X_1 , X_2 , and X_3 could be absent from the mutant (i.e., no amino acid corresponding to the position). The most preferred peroxisome-targeting-avoiding sequence is IAV, where V is at the carboxy-terminus.

35 In another of its aspects, the invention entails a combination of luciferases, in a cell (eukaryotic or prokaryotic), a solution (free or linked as a reporter to

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an antibody, antibody-fragment, nucleic acid probe, or the like), or adhered to a solid surface, optionally through an antibody, antibody fragment or nucleic acid, and exposed to a solution, provided that at least one of the luciferases is a mutant, both of the luciferases remain active in producing bioluminescence, and the wavelengths of peak intensities of the bioluminescence of the luciferases differ because the amino acid sequences of the luciferases differ at at least one of the positions corresponding to positions 214, 215, 223, 224, 232, 236, 237, 238, 242, 244, 245, 247, 248, 282, 283 and 348 in the amino acid sequence of LucPplGR, provided that one or both of the luciferases optionally have peroxisome-targeting-avoiding sequences.

In another of its aspects, the invention entails a DNA molecule, which may be an eukaryotic or prokaryotic expression vector, which comprises a segment which has a sequence which encodes a mutant beetle luciferase of the invention.

Most preferred among the DNAs of the invention are those with segments which encode a preferred mutant luciferase of the invention.

From the description of the invention provided herein, the skilled will recognize many modifications and variations of what has been described that are within the spirit of the invention. It is intended that such modifications and variations also be understood as part of the invention.

SEQUENCE LISTING

(1) GENERAL INFORMATION:

- (i) APPLICANT: Promega Corporation
(ii) TITLE OF INVENTION: Mutant Luciferases
(iii) NUMBER OF SEQUENCES: 1
(iv) CORRESPONDENCE ADDRESS:
 (A) ADDRESSEE: Foley & Lardner
 (B) STREET: P. O. Box 1497
 (C) CITY: Madison
 (D) STATE: Wisconsin
 (E) COUNTRY: US
 (F) ZIP: 53701-1497
(vii) PRIOR APPLICATION DATA:
 (A) APPLICATION NUMBER: US 08/177,081
 (B) FILING DATE: 3-Jan-1994
(viii) ATTORNEY/AGENT INFORMATION:
 (A) NAME: Scanlon, William J.
 (B) REGISTRATION NUMBER: 30136
 (C) REFERENCE/DOCKET NUMBER: 19017/148P
(ix) TELECOMMUNICATION INFORMATION:
 (A) TELEPHONE: (608) 258-4284
 (B) TELEFAX: (608) 258-4258

(2) INFORMATION FOR SEQ ID NO:1:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1632 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear
 (ii) MOLECULE TYPE: cDNA to mRNA
 (iii) HYPOTHETICAL: no
 (iv) ANTISENSE: no
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

ATG Met	ATG Met	AAG Lys	AGA Arg	GAG Glu 5	AAA Lys	AAT Asn	GTT Val	GTA Val	TAT Tyr 10	GGA Gly	CCC Pro	GAA Glu	CCC Pro	CTA Leu 15	CAC His	48
CCC Pro	TTG Leu	GAA Glu	GAC Asp 20	TTA Leu	ACA Thr	GCA Ala	GGA Gly	GAA Glu 25	ATG Met	CTC Leu	TTC Phe	AGG Arg	GCC Ala 30	CTT Leu	CGA Arg	96
AAA Lys	CAT His	TCT Ser 35	CAT His	TTA Leu	CCG Pro	CAG Gln	GCT Ala 40	TTA Leu	GTA Val	GAT Asp	GTG Val	TAT Tyr 45	GGT Gly	GAA Glu	GAA Glu	144
TGG Trp	ATT Ile 50	TCA Ser	TAT Tyr	AAA Lys	GAG Glu	TTT Phe 55	TTT Phe	GAA Glu	ACT Thr	ACA Thr	TGC Cys 60	CTA Leu	CTA Leu	GCA Ala	CAA Gln	192
AGT Ser 65	CTT Leu	CAC His	AAT Asn	TGT Cys	GCA Gly 70	TAC Tyr	AAG Lys	ATG Met	AGT Ser	GAT Asp 75	GTA Val	GTG Val	TCG Ser	ATC Ile	TGC Cys 80	240
GCG Ala	GAG Glu	AAC Asn	AAT Asn	AAA Lys 85	AGA Arg	TTT Phe	TTT Phe	GTT Val	CCC Pro 90	ATT Ile	ATT Ile	GCA Ala	GCT Ala	TGG Trp 95	TAT Tyr	288
ATT Ile	GGT Gly	ATG Met	ATT Ile 100	GTA Val	GCA Ala	CCT Pro	GTT Val	AAT Asn 105	GAG Glu	GGC Gly	TAC Tyr	ATC Ile	CCA Pro 110	GAT Asp	GAA Glu	336
CTC Leu	TGT Cys	AAG Lys 115	GTC Val	ATG Met	GGT Gly	ATA Ile	TCG Ser 120	AGA Arg	CCA Pro	CAA Gln	CTA Leu	GTT Val 125	TTT Phe	TGT Cys	ACA Thr	384

AAG AAT ATT CTA AAT AAG GTA TTG GAG GTA CAG AGC AGA ACT GAT TTC Lys Asn Ile Leu Asn Lys Val Leu Glu Val Gln Ser Arg Thr Asp Phe 130 135 140	432
ATA AAA AGG ATT ATC ATA CTA GAT GCT GTA GAA AAC ATA CAC GGT TGT Ile Lys Arg Ile Ile Ile Leu Asp Ala Val Glu Asn Ile His Gly Cys 145 150 155 160	480
GAA AGT CTT CCC AAT TTT ATT TCT CGT TAT TCG GAT GGA AAT ATT GCC Glu Ser Leu Pro Asn Phe Ile Ser Arg Tyr Ser Asp Gly Asn Ile Ala 165 170 175	528
AAC TTC AAA CCT TTA CAT TAC GAT CCT GTT GAA CAA GTG GCA GCT ATC Asn Phe Lys Pro Leu His Tyr Asp Pro Val Glu Gln Val Ala Ala Ile 180 185 190	576
TTA TGT TCG TCA GGC ACA ACT GGA TTA CCG AAA GGT GTA ATG CAA ACT Leu Cys Ser Ser Gly Thr Thr Gly Leu Pro Lys Gly Val Met Gln Thr 195 200 205	624
CAT AGA AAT GTT TGT GTC CGA CTT ATA CAT GCT TTA GAC CCC AGG GTA His Arg Asn Val Cys Val Arg Leu Ile His Ala Leu Asp Pro Arg Val 210 215 220	672
GGA ACG CAA CTT ATT CCT GGT GTG ACA GTC TTA GTA TAT CTG CCT TTT Gly Thr Gln Leu Ile Pro Gly Val Thr Val Leu Val Tyr Leu Pro Phe 225 230 235 240	720
TTC CAT GCT TTT GGG TTC TCT ATA AAC TTG GGA TAC TTC ATG GTG GGT Phe His Ala Phe Gly Phe Ser Ile Asn Leu Gly Tyr Phe Met Val Gly 245 250 255	768
CTT CGT GTT ATC ATG TTA AGA CGA TTT GAT CAA GAA GCA TTT CTA AAA Leu Arg Val Ile Met Leu Arg Arg Phe Asp Gln Glu Ala Phe Leu Lys 260 265 270	816
GCT ATT CAG GAT TAT GAA GTT CGA AGT GTA ATT AAC GTT CCA GCA ATA Ala Ile Gln Asp Tyr Glu Val Arg Ser Val Ile Asn Val Pro Ala Ile 275 280 285	864
ATA TTG TTC TTA TCG AAA AGT CCT TTG GTT GAC AAA TAC GAT TTA TCA Ile Leu Phe Leu Ser Lys Ser Pro Leu Val Asp Lys Tyr Asp Leu Ser 290 295 300	912
AGT TTA AGG GAA TTG TGT TGC GGT GCG GCA CCA TTA GCA AAC GAA GTT Ser Leu Arg Glu Leu Cys Cys Gly Ala Ala Pro Leu Ala Lys Glu Val 305 310 315 320	960
GCT GAG ATT GCA GTA AAA CGA TTA AAC TTG CCA GGA ATT CGC TGT GGA Ala Glu Ile Ala Val Lys Arg Leu Asn Leu Pro Gly Ile Arg Cys Gly 325 330 335	1008
TTT GGT TTG ACA GAA TCT ACT TCA GCT AAT ATA CAC AGT CTT AGG GAT Phe Gly Leu Thr Glu Ser Thr Ser Ala Asn Ile His Ser Leu Arg Asp 340 345 350	1056
GAA TTT AAA TCA GGA TCA CTT GGA AGA GTT ACT CCT TTA ATG GCA GCT Glu Phe Lys Ser Gly Ser Leu Gly Arg Val Thr Pro Leu Met Ala Ala 355 360 365	1104
AAA ATA GCA GAT AGG GAA ACT GGT AAA GCA TTG GGA CCA AAT CAA GTT Lys Ile Ala Asp Arg Glu Thr Gly Lys Ala Leu Gly Pro Asn Gln Val 370 375 380	1152
GGT GAA TTA TGC ATT AAA GGT CCC ATG GTA TCG AAA GGT TAC GTG AAC Gly Glu Leu Cys Ile Lys Gly Pro Met Val Ser Lys Gly Tyr Val Asn 385 390 395 400	1200

AAT GTA GAA GCT ACC AAA GAA GCT ATT GAT GAT GAT GGT TGG CTT CAC	1248
Asn Val Glu Ala Thr Lys Glu Ala Ile Asp Asp Asp Gly Trp Leu His	
405 410 415	
TCT GGA GAC TTT GGA TAC TAT GAT GAG GAT GAG CAT TTC TAT GTG GTG	1296
Ser Gly Asp Phe Gly Tyr Tyr Asp Glu Asp Glu His Phe Tyr Val Val	
420 425 430	
GAC CGT TAC AAG GAA TTG ATT AAA TAT AAG GGC TCT CAG GTA GCA CCT	1344
Asp Arg Tyr Lys Glu Leu Ile Lys Tyr Lys Gly Ser Gln Val Ala Pro	
435 440 445	
GCA GAA CTA GAA GAG ATT TTA TTG AAA AAT CCA TGT ATC AGA GAT GTT	1392
Ala Glu Leu Glu Glu Ile Leu Leu Lys Asn Pro Cys Ile Arg Asp Val	
450 455 460	
GCT GTG GTT GGT ATT CCT GAT CTA GAA GCT GGA GAA CTG CCA TCT GCG	1440
Ala Val Val Gly Ile Pro Asp Leu Glu Ala Gly Glu Leu Pro Ser Ala	
465 470 475 480	
TTT GTG GTT ATA CAG CCC GGA AAG GAG ATT ACA GCT AAA GAA GTT TAC	1488
Phe Val Val Ile Gln Pro Gly Lys Glu Ile Thr Ala Lys Glu Val Tyr	
485 490 495	
GAT TAT CTT GCC GAG AGG GTC TCC CAT ACA AAG TAT TTG CGT GGA GGG	1536
Asp Tyr Leu Ala Glu Arg Val Ser His Thr Lys Tyr Leu Arg Gly Gly	
500 505 510	
GTT CGA TTC GTT GAT AGC ATA CCA AGG AAT GTT ACA GGT AAA ATT ACA	1584
Val Arg Phe Val Asp Ser Ile Pro Arg Asn Val Thr Gly Lys Ile Thr	
515 520 525	
AGA AAG GAA CTT CTG AAG CAG TTG CTG GAG AAG AGT TCT AAA CTT TAA	1632
Arg Lys Glu Leu Leu Lys Gln Leu Leu Glu Lys Ser Ser Lys Leu	
530 535 540	

CLAIMS

1. A mutant beetle luciferase which has an amino acid sequence that differs from that of the corresponding wild-type luciferase by a substitution at one position or
5 substitutions at two positions; provided that, if there is a substitution at one position, the position corresponds to a position in the amino acid sequence of LucPplGR selected from the group consisting of position
10 214, 215, 223, 224, 232, 236, 237, 238, 242, 244, 245, 247, 248, 282, 283 and 348; and provided further that, if there are substitutions at two positions, at least one of the positions corresponds to a position in the amino acid sequence of LucPplGR selected from the group consisting
15 of position 214, 215, 223, 224, 232, 236, 237, 238, 242, 244, 245, 247, 248, 282, 283 and 348.
2. A mutant luciferase according to Claim 1 wherein there is one amino acid substitution.
3. A mutant luciferase according to Claim 1 wherein there are two amino acid substitutions.
- 20 4. A mutant luciferase according to Claim 3 wherein each of the amino acid substitutions is at a position corresponding to a position in the amino acid sequence of LucPplGR selected from the group consisting of position 214, 215, 223, 224, 232, 236, 237, 238, 242,
25 244, 245, 247, 248, 282, 283 and 348.
5. A mutant luciferase according to Claim 1 wherein the corresponding wild-type luciferase is selected from the group consisting of LucPplGR, LucPplyG, LucPplyE, LucPplOR, the luciferase of *Photinus pyralis*,
30 the luciferase of *Luciola cruciata*, the luciferase of *Luciola lateralis*, and the luciferase of *Luciola mingrelica*.

6. A mutant luciferase according to Claim 2 wherein the corresponding wild-type luciferase is selected from the group consisting of LucPplGR, LucPplyG, LucPplyE, LucPplOR, the luciferase of *Photinus pyralis*,
5 the luciferase of *Luciola cruciata*, the luciferase of *Luciola lateralis*, and the luciferase of *Luciola mingrelica*.

7. A mutant luciferase according to Claim 3 wherein the corresponding wild-type luciferase is
10 selected from the group consisting of LucPplGR, LucPplyG, LucPplyE, LucPplOR, the luciferase of *Photinus pyralis*, the luciferase of *Luciola cruciata*, the luciferase of *Luciola lateralis*, and the luciferase of *Luciola mingrelica*.

15 8. A mutant luciferase according to Claim 4 wherein the corresponding wild-type luciferase is selected from the group consisting of LucPplGR, LucPplyG, LucPplyE, LucPplOR, the luciferase of *Photinus pyralis*, the luciferase of *Luciola cruciata*, the luciferase of
20 *Luciola lateralis*, and the luciferase of *Luciola mingrelica*.

9. A mutant luciferase according to Claim 5 wherein the corresponding wild-type luciferase is selected from the group consisting of LucPplGR, LucPplyG,
25 LucPplyE, and LucPplOR.

10. A mutant luciferase according to Claim 6 wherein the corresponding wild-type luciferase is selected from the group consisting of LucPplGR, LucPplyG, LucPplyE, and LucPplOR.

30 11. A mutant luciferase according to Claim 7 wherein the corresponding wild-type luciferase is selected from the group consisting of LucPplGR, LucPplyG, LucPplyE, and LucPplOR.

35 12. A mutant luciferase according to Claim 8 wherein the corresponding wild-type luciferase is selected from the group consisting of LucPplGR, LucPplyG, LucPplyE, and LucPplOR.

13. A mutant luciferase of Claim 9 wherein the corresponding wild-type luciferase is LucPplGR.

14. A mutant luciferase of Claim 10 wherein the corresponding wild-type luciferase is LucPplGR.

5 15. A mutant luciferase of Claim 11 wherein the corresponding wild-type luciferase is LucPplGR.

16. A mutant luciferase of Claim 12 wherein the corresponding wild-type luciferase is LucPplGR.

10 17. A mutant luciferase of Claim 13 wherein the mutant is selected from the group consisting of
LucPplGR-R₂₁₅H, -R₂₁₅G, -R₂₁₅T, -R₂₁₅M, -R₂₁₅P, -R₂₁₅A, -R₂₁₅L,
-R₂₂₃L, -R₂₂₃Q, -R₂₂₃M, -R₂₂₃H, -V₂₂₄I, -V₂₂₄S, -V₂₂₄F, -V₂₂₄Y, -V₂₂₄L,
-V₂₂₄H, -V₂₂₄G, -V₂₃₂E, -V₂₃₆H, -V₂₃₆W, -Y₂₃₇S, -Y₂₃₇C, -L₂₃₈R, -L₂₃₈M,
-L₂₃₈Q, -L₂₃₈S, -L₂₃₈D, -H₂₄₂A, -F₂₄₄L, -G₂₄₅S, -G₂₄₅E, -S₂₄₇H, -S₂₄₇T,
15 -S₂₄₇Y, -S₂₄₇F, -I₂₄₈R, -I₂₄₈V, -I₂₄₈F, -I₂₄₈T, -I₂₄₈S, -I₂₄₈N, -H₃₄₈N,
-H₃₄₈Q, -H₃₄₈E, -H₃₄₈C, -S₂₄₇F/F₂₄₆L, -S₂₄₇F/I₂₄₈C, -S₂₄₇F/I₂₄₈T,
-V₂₂₄F/R₂₁₅G, -V₂₂₄F/R₂₁₅T, -V₂₂₄F/R₂₁₅V, -V₂₂₄F/R₂₁₅P, -V₂₂₄F/P₂₂₂S,
-V₂₂₄F/Q₂₂₇E, -V₂₂₄F/L₂₃₈V, -V₂₂₄F/L₂₃₈T, -V₂₂₄F/S₂₄₇G, -V₂₂₄F/S₂₄₇H,
-V₂₂₄F/S₂₄₇T, and -V₂₂₄F/S₂₄₇F.

20 18. A DNA molecule which comprises a segment which has a sequence which encodes a mutant beetle luciferase which has an amino acid sequence that differs from that of the corresponding wild-type luciferase by a
substitution at one position or substitutions at two
25 positions; provided that, if there is a substitution at one position, the position corresponds to a position in the amino acid sequence of LucPplGR selected from the group consisting of position 214, 215, 223, 224, 232, 236, 237, 238, 242, 244, 245, 247, 248, 282, 283 and 348;
30 and provided further that, if there are substitutions at two positions, at least one of the positions corresponds to a position in the amino acid sequence of LucPplGR selected from the group consisting of position 214, 215, 223, 224, 232, 236, 237, 238, 242, 244, 245, 247, 248,
35 282, 283 and 348.

19. A DNA molecule according to Claim 18 wherein the encoded mutant luciferase has one amino acid substitution.

20. A DNA molecule according to Claim 18 wherein
5 the encoded mutant luciferase has two amino acid substitutions.

21. A DNA molecule according to Claim 20 wherein, in the encoded mutant luciferase, each of the amino acid substitutions is at a position corresponding to a
10 position in the amino acid sequence of LucPplGR selected from the group consisting of position 214, 215, 223, 224, 232, 236, 237, 238, 242, 244, 245, 247, 248, 282, 283 and 348.

22. A DNA molecule according to Claim 18 wherein,
15 for the encoded amino acid sequence, the corresponding wild-type luciferase is selected from the group consisting of LucPplGR, LucPplyG, LucPplyE, LucPplOR, the luciferase of *Photinus pyralis*, the luciferase of *Luciola cruciata*, the luciferase of *Luciola lateralis*, and the
20 luciferase of *Luciola mingrelica*.

23. A DNA molecule according to Claim 19 wherein, for the encoded amino acid sequence, the corresponding wild-type luciferase is selected from the group consisting of LucPplGR, LucPplyG, LucPplyE, LucPplOR, the
25 luciferase of *Photinus pyralis*, the luciferase of *Luciola cruciata*, the luciferase of *Luciola lateralis*, and the luciferase of *Luciola mingrelica*.

24. A DNA molecule according to Claim 20 wherein, for the encoded amino acid sequence, the corresponding
30 wild-type luciferase is selected from the group consisting of LucPplGR, LucPplyG, LucPplyE, LucPplOR, the luciferase of *Photinus pyralis*, the luciferase of *Luciola cruciata*, the luciferase of *Luciola lateralis*, and the luciferase of *Luciola mingrelica*.

25. A DNA molecule according to Claim 21 wherein,
for the encoded amino acid sequence, the corresponding
wild-type luciferase is selected from the group
consisting of LucPplGR, LucPplyG, LucPplyE, LucPplOR, the
5 luciferase of *Photinus pyralis*, the luciferase of *Luciola*
cruciata, the luciferase of *Luciola lateralis*, and the
luciferase of *Luciola mingrelica*.

26. A DNA molecule according to Claim 22 wherein,
for the encoded amino acid sequence, the corresponding
10 wild-type luciferase is selected from the group
consisting of LucPplGR, LucPplyG, LucPplyE, and LucPplOR.

27. A mutant luciferase according to Claim 23
wherein, for the encoded amino acid sequence, the
corresponding wild-type luciferase is selected from the
15 group consisting of LucPplGR, LucPplyG, LucPplyE, and
LucPplOR.

28. A DNA molecule according to Claim 24 wherein,
for the encoded amino acid sequence, the corresponding
wild-type luciferase is selected from the group
20 consisting of LucPplGR, LucPplyG, LucPplyE, and LucPplOR.

29. A DNA molecule according to Claim 25 wherein,
for the encoded amino acid sequence, the corresponding
wild-type luciferase is selected from the group
consisting of LucPplGR, LucPplyG, LucPplyE, and LucPplOR.

30. A DNA molecule according to Claim 26 wherein,
for the encoded amino acid sequence, the corresponding
wild-type luciferase is LucPplGR.

31. A DNA molecule according to Claim 27 wherein,
for the encoded amino acid sequence, the corresponding
30 wild-type luciferase is LucPplGR.

32. A DNA molecule according to Claim 28 wherein,
for the encoded amino acid sequence, the corresponding
wild-type luciferase is LucPplGR.

33. A DNA molecule according to Claim 29 wherein,
for the encoded amino acid sequence, the corresponding
35 wild-type luciferase is LucPplGR.

34. A DNA molecule according to Claim 30 wherein the encoded mutant luciferase is selected from the group consisting of LucPplGR-R₂₁₅H, -R₂₁₅G, -R₂₁₅T, -R₂₁₅M, -R₂₁₅P, -R₂₁₅A, -R₂₁₅L, -R₂₂₃L, -R₂₂₃Q, -R₂₂₃M, -R₂₂₃H, -V₂₂₄I, -V₂₂₄S, -V₂₂₄F, -V₂₂₄Y, -V₂₂₄L, -V₂₂₄H, -V₂₂₄G, -V₂₃₂E, -V₂₃₆H, -V₂₃₆W, -Y₂₃₇S, -Y₂₃₇C, -L₂₃₈R, -L₂₃₈M, -L₂₃₈Q, -L₂₃₈S, -L₂₃₈D, -H₂₄₂A, -F₂₄₄L, -G₂₄₅S, -G₂₄₅E, -S₂₄₇H, -S₂₄₇T, -S₂₄₇Y, -S₂₄₇F, -I₂₄₈R, -I₂₄₈V, -I₂₄₈F, -I₂₄₈T, -I₂₄₈S, -I₂₄₈N, -H₃₄₈N, -H₃₄₈Q, -H₃₄₈E, -H₃₄₈C, -S₂₄₇F/F₂₄₆L, -S₂₄₇F/I₂₄₈C, -S₂₄₇F/I₂₄₈T, -V₂₂₄F/R₂₁₅G, -V₂₂₄F/R₂₁₅T, -V₂₂₄F/R₂₁₅V, -V₂₂₄F/R₂₁₅P, -V₂₂₄F/P₂₂₂S, -V₂₂₄F/Q₂₂₇E, -V₂₂₄F/L₂₃₈V, -V₂₂₄F/L₂₃₈T, -V₂₂₄F/S₂₄₇G, -V₂₂₄F/S₂₄₇H, -V₂₂₄F/S₂₄₇T, and -V₂₂₄F/S₂₄₇F.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US95/00108

A. CLASSIFICATION OF SUBJECT MATTER IPC(6) : C12N 9/02, 15/53 US CL : 435/189; 536/23.2 According to International Patent Classification (IPC) or to both national classification and IPC				
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) U.S. : 435/189, 172.3, 320.1, 252.3, 252.33; 536/23.2 Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) Please See Extra Sheet.				
C. DOCUMENTS CONSIDERED TO BE RELEVANT				
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.		
X	FEBS Letters, Volume 307, No. 2, issued July 1992, G. Sala-Newby et al., "Engineering Firefly Luciferase as an Indicator of Cyclic AMP-Dependent Protein Kinase in Living Cells", pages 241-244, see entire document.	1, 2, 5, 6, 18, 19, 22, 23		
X	BIOCHEMICAL JOURNAL, Volume 279, issued November 1991, G. Sala-Newby et al., "Engineering a Bioluminescent Indicator for Cyclic AMP-Dependent Protein Kinase", pages 727-732, see entire document.	1, 2, 5, 6, 18, 19, 22, 23		
A	PROTEIN ENGINEERING, Volume 4, No. 6, issued August 1991, N. Kajiyama et al., "Isolation and Characterization of Mutants of Firefly Luciferase Which Produce Different Colors of Light", pages 691-693.	1-34		
<input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C. <input type="checkbox"/> See patent family annex.				
<table border="0"> <tr> <td> * Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed </td> <td> "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "Z" document member of the same patent family </td> </tr> </table>			* Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "Z" document member of the same patent family
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Date of the actual completion of the international search 05 APRIL 1995		Date of mailing of the international search report 17 APR 1995		
Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Facsimile No. (703) 305-3230		Authorized officer <i>Rebecca Prouty</i> REBECCA PROUTY Telephone No. (703) 308-0196		

INTERNATIONAL SEARCH REPORTInter. .onal application No.
PCT/US95/00108**C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	JOURNAL OF BIOLUMINESCENCE AND CHEMILUMINESCENCE, Volume 5, issued April 1990, K.V. Wood, "Luc Genes: Introduction of Colour Into Bioluminescence Assays", pages 107-114.	1-34
A	JOURNAL OF BIOLUMINESCENCE AND CHEMILUMINESCENCE, Volume 4, issued July 1989, K.V. Wood et al., "Introduction to Beetle Luciferases and Their Applications", pages 289-301.	1-34
A	JOURNAL OF BIOLUMINESCENCE AND CHEMILUMINESCENCE, Volume 4, issued July 1989, K.V. Wood et al., "Bioluminescent Click Beetles Revisited", pages 31-39.	1-34
A	SCIENCE, Volume 244, issued 12 May 1989, K.V. Wood et al., "Complementary DNA Coding Click Betle Luciferases Can Elicit Bioluminescence of Different Colors", pages 700-702.	1-34

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International application No.
PCT/US95/00108

B. FIELDS SEARCHED

Electronic data bases consulted (Name of data base and where practicable terms used):

APS, MEDLINE, BIOSIS, LIFESCI, EMBASE, WPI, BIOTECHDS, CA

search terms: luciferase#, muta? or modif?, gene# or sequence#, beetle# or firefl?, pyrophorus or plagiophthalmus,
photinus or luciola

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